

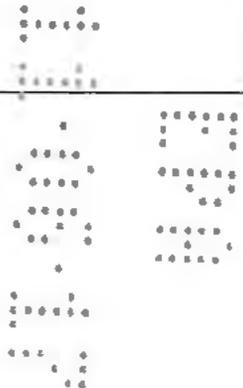
Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information

-001

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area. Page # 1 of 3

Row 1 Administrative Data	Reporter name: [REDACTED]	Submission date:	Contact person (if different than reporter)	Internal ID 1-49655563
	Address: Ontario		Address:	
	Phone #: [REDACTED]		Phone #:	
	Incident Status: New	Location and date of incident Ontario 08/27/2017	Date registrant became aware of incident: 9/10/2017	Was incident part of larger study?
Row 2 Pesticide(s) Involved	EPA Registration # (Product 1) 24359	EPA Registration # (Product 2)	EPA Registration # (Product 3)	
	A.I. (s)	A.I. (s)	A.I. (s)	
	Product 1 Name Glyphos Soluble Concentrate Herbicide Canada	Product 2 Name	Product 3 Name	
	Exposed to concentrate prior to dilution? NA	Exposed to concentrate prior to dilution?	Exposed to concentrate prior to dilution?	
	Formulation	Formulation	Formulation	
Row 3 Incident Circumstances	Evidence label directions were not followed? No Intentional misuse? No	Incident site: (examples include home, yard, school, industrial, nursery/greenhouse, surface water, commercial turf, building/office, forest/ woods, agricultural (specify crop) right-of-way (rail, utility, highway))  Own Residence	Situation: (act of using product): (examples include mixing/loading, reentry, application, transportation, repair/maintenance of application equipment, manufacturing/ formulating)  See Description Notes	
	Applicator certified PCO? Not applicable			
	How exposed: (examples include direct contact with treated surface, ingestion, spill, drift, runoff)  See Incident Description			

\*Personal privacy information\*



*9/10/2017 10:05:58 AM Glyphos Soluble Concentrate Herbicide Canada  
PCP #24359*

*HX: The caller got some of the product on his hand 2 weeks ago. He did take a shower after exposure. Shortly after exposure he started to experience joint and muscle pain. Could his symptoms be related to this product?*

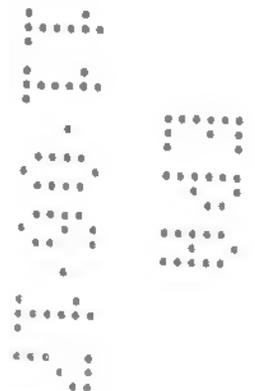
*A: The symptoms described would not be expected from exposure to this product.*

- Product may cause temporary skin irritation.*
- Consider other causes for your symptoms.*
- Please call back with any further questions or concerns.*

*Consulted with AB*

*10/6/2017 11:48:10 AM PMRA report generated.*

*10/11/2017 9:44:57 PM PMRA report sent.*



**Voluntary Industry Reporting Form for 6(a)(2) Incident Information Involving Humans**

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area. Page # 3 of 3

Demographic information Age: <i>Unknown</i> Sex: <i>Male</i> Occupation: (if relevant)	Exposure route: <i>Dermal</i>	Was adverse effect result of suicide/homicide or attempted suicide/homicide? <i>No</i>	Was protective clothing worn (specify)?  <i>Not applicable</i>
If female, pregnant? <i>Did not query</i>	Was exposure occupational? <i>No</i> If yes, days lost due to illness:	Time between exposure and onset of symptoms: <i>See Symptoms</i>	
Type of medical care sought: (examples include none, clinic, hospital emergency department, private physician, PCC, hospital inpatient). <i>On-site</i>	List signs/symptoms/adverse effects.  <i>Joint pain, 24 hrs or less;</i> <i>Other miscellaneous - Muscle pain, 24 hrs or less;</i>		If lab tests were performed, list test names and results (if available, submit reports).  <i>Not Reported</i>
Exposure data: Amount of pesticide: Exposure duration: Weight:			
Human severity category: <i>HC</i>			

This box can be used to provide any explanatory or qualifying information surrounding the incident. (add additional pages if necessary)

Internal ID #  
*1-49655563*